

(12) United States Patent

Ballesteros et al.

(10) **Patent No.:**

US 9,168,394 B2

(45) **Date of Patent:**

*Oct. 27, 2015

(54)	PIGMEN	TED SKIN	-CARE COMPOSITIONS		5,741,355 5,753,026		4/1998 5/1998	Yamamoto Kuntz
(71)	Annlicant:	Johnson &	& Johnson Consumer		5,776,497		7/1998	Lagrange
(11)	rippiicani.		es, Inc., Skillman, NJ (US)		5,807,497			Gailberger
		Companie	25, The., Skinman, NJ (03)		5,851,277			Muller Rees
(72)	Inventors	Ann Theo	dore Ballesteros, New		5,958,125			Schmid
(12)	mvemors.		t, NJ (US); Stephen William		5,972,098 6,001,373		10/1999	Igo Kemenes
			ingdon Valley, PA (US)		6,113,683		9/2000	
		riu, muni	ingdon variey, FA (OS)		6,117,435		9/2000	
(73)	Assignee.	Johnson &	& Johnson Consumer Inc.,		6,132,504		10/2000	Kuntz
(13)	1 issignee.	Skillman,			6,132,873		10/2000	
		ommun,	(05)		6,156,115		12/2000	
(*)	Notice:	Subject to	any disclaimer, the term of th	nis	6,187,298 6,190,445	BI	2/2001	Noguchi
` /			extended or adjusted under		6,190,648		2/2001	
			l(b) by 0 days.		6,238,471		5/2001	
			• •		6,238,472		5/2001	
			it is subject to a terminal di	IS-	6,267,810		7/2001	
		claimer.			6,280,520		8/2001	
(21)	Anni No	12/700 097	7		6,306,409		10/2001	
(21)	Appi. No	13/799,087	,		6,428,773 6,451,294		8/2002 9/2002	
(22)	Filed:	Mar. 13, 2	013		6,475,500		11/2002	
(22)	i nea.	WIGH. 13, 2	013		6,485,556			DeLuca, Jr.
(65)		Prior P	ublication Data		6,500,251		12/2002	
()	770 004 4/0				6,508,876			Bernhardt
	US 2014/0	271740 A1	Sep. 18, 2014		6,511,672		1/2003	
(51)	T 4 63				6,517,628 6,524,598		2/2003 2/2003	
(51)	Int. Cl.		(2007, 01)		6,531,221			Schuhmacher
	A61K 8/02		(2006.01)		6,545,809			Phillips
(50)	A61Q 1/02	•	(2006.01)		6,579,355	B1		Schmidt
(52)	U.S. Cl.	4676	0 1/03 (2012 01): 4/1W 0/03	50	6,596,070			Schmidt
			Q 1/02 (2013.01); A61K 8/02.		6,599,355			Schmidt
	(2		61K 2800/412 (2013.01); A61		6,630,018 6,632,275		10/2003 10/2003	
		2800/4.	36 (2013.01); A61K 2800/59.		6,641,823	B2	11/2003	
(50)	E: 11 . CC	1	(2013.0	1)	6,641,874		11/2003	
(58)		lassification		00			(Con	tinued)
			A61Q 1/0	02			(COII	imucu)
	See applica	ation me ioi	r complete search history.		FO	REIG	N PATE	NT DOCUMENTS
(56)		Referen	ices Cited					
(00)				EP			9109 B1	2/1977
	U.	S. PATENT	DOCUMENTS	EP EP			5889 A3 5523 B1	9/1984 1/1991
		0/40=0	m.t	EP			7747 B1	5/1992
	3,692,768 A 3,978,207 A	9/1972 8/1976	Takata	EP			1047 B1	8/1994
	4,456,486 A		Bernhard	EP			5325 B1	11/1994
	4,457,784 A		Bernhard	EP EP			2329 B1	7/1995
	4,494,993 A		Bernhard	EP			0022 B1 2110 B1	1/1996 9/1996
	4,552,593 A 4,565,581 A		Ostertag Bernhard	EP			5486 B1	8/1997
	4,623,396 A		Kimura				(Con	tinued)
	4,648,908 A		Takasuka				(COII	imaca)
	4,710,375 A		Takasuka		-	_		
	4,828,826 A	5/1989 8/1990		Prin	nary Exam	iner –	— Heana .	Popa
-	4,952,245 A 5,116,664 A		Kimura	Assi	stant Exan	iner-	— Nicole	Babson
	5,156,678 A		Glausch					
:	5,169,442 A		Noguchi					
	5,302,199 A		Prengel	(57)			ABST	FRACT
	5,326,392 A 5,364,467 A	7/1994 11/1994	Schmid		. 1 1			
	5,496,543 A	3/1996	Lagrange					sition including a dermatologi-
;	5,607,504 A	3/1997	Schmid					at least a first and second inter-
	5,618,342 A		Herget					ition according to the invention
	5,624,486 A 5,624,487 A		Schmid Schmidt					appearance of the skin and in
	5,624,731 A		Desjardins	part	icular in re	ducin	g the app	earance of skin discontinuities.
:	5,662,738 A	9/1997	Schmid					
	5,690,916 A		Kimura			12	Claima	No Drawings
	5,733,658 A	5/1998	Schmid			13	Ciaiiis,	No Drawings

US 9,168,394 B2Page 2

(56)			Referen	ces Cited	8,083,846			Zimmermann
	1	101	DATENIT	DOCUMENTS	8,088,212 8,088,214		1/2012	Bagala, Sr. Fuller
	,	J.S. 1	ALENI	DOCUMENTS	8,114,211			Handrosch
6.6	48,957	B1	11/2003	Andes	8,114,388		2/2012	Simon
	56,259		12/2003		8,129,021		3/2012	
	63,852		12/2003	Simon	8,147,853		4/2012	Taylor
	89,205			Brückner	8,158,566 2001/0001174		4/2012 5/2001	Wei Andes
	92,561		2/2004		2002/001174		2/2002	
	96,049 06,109		2/2004 3/2004	DeLuca, Jr.	2002/0018791		2/2002	
	19,837		4/2004		2002/0033117	A1	3/2002	Inoue
	19,838		4/2004		2002/0064509		5/2002	
,	43,285			Anselmann	2002/0134282			Ostertag
	44,559		6/2004		2002/0169244 2003/0017124			Ostertag Agostini
	47,073 49,936		6/2004	Argoitia	2003/0017124			Poetsch
	51,022		6/2004		2003/0064039			Kolodziej
	73,499		8/2004		2003/0075079			Sommer
	83,584			Takahashi	2003/0091813		5/2003	
	94,037			Zimmermann	2003/0118622 2003/0157041		6/2003 8/2003	
	00,125			Zimmermann	2003/0157041		8/2003	
	21,333 31,785		12/2004	Zimmermann	2003/0209169		11/2003	
	33,959		12/2004		2003/0211058	A1		Matts
	37,925		1/2005		2004/0052743			Schmidt
	40,993			Schmidt	2004/0057915		3/2004 4/2004	Gers Barlag
	75,264			Zimmermann	2004/0076650 2004/0105827		6/2004	Blin Grimm
	84,289 06,015		4/2005	Schoen Shiloach	2004/0177788		9/2004	Rick
	14,700			DeLuca, Jr.	2004/0191198		9/2004	Hochstein
	45,007			Bagala, Sr.	2004/0194663		10/2004	Li
	60,126		6/2006	Andes	2004/0219116		11/2004	Reynders
	77,897			Brueckner	2004/0223991 2004/0223992		11/2004 11/2004	Wei
	69,223		1/2007		2004/0223992			Clapp
	02,199 26,503			Shiloach Anselmann	2004/0234564		11/2004	
	41,503			Noguchi	2004/0234565		11/2004	
	55,736		8/2007		2004/0241118		12/2004	
	64,670		9/2007		2004/0258640 2005/0001203		1/2004	Simon Bertaux
	00,510 03,622		11/2007 12/2007		2005/0001205			Duffournier et al.
	18,861			Bagala, Sr.	2005/0061205	A1		Kobayashi
	26,671			Shiloach	2005/0112072		5/2005	Wang
	44,590			Schmidt	2005/0142084 2005/0143269		6/2005 6/2005	Ganguly Wei
	65,109 87,669		4/2008 6/2008	Rathschlag	2005/0143205			Dabkowski
	96,401			Jungnitz	2005/0175562			Hadasch
	52,597		11/2008		2005/0176850		8/2005	Schmidt
	55,726			Schoenefeld	2005/0204958			Kuebelbeck
	79,323			Rathschlag	2005/0220735 2005/0220736		10/2005 10/2005	
	85,183 17,404		4/2009	Hochstein Buiged	2005/0252410		11/2005	
	31,184		5/2009	· .	2005/0268405	A1	12/2005	
	78,879		8/2009	Huber	2005/0273947		12/2005	
	79,079		8/2009		2005/0276768 2005/0276779		12/2005 12/2005	
	04,862 21,966		10/2009	Ambrosius	2006/0005742			Moeschl
	28,998		12/2009		2006/0013838		1/2006	Peng
	82,604		3/2010		2006/0027140		2/2006	
,	91,196		4/2010		2006/0032404		2/2006	
	08,823		5/2010		2006/0034787 2006/0047018		2/2006 3/2006	
	45,003 67,214		8/2010	Hennemann Simon	2006/0051204		3/2006	
	72,214		8/2010		2006/0051304		3/2006	
	80,955		8/2010		2006/0088483		4/2006	Thevenet
	94,740		9/2010		2006/0144294 2006/0147390		7/2006 7/2006	Misaki Schreiber
	99,746 20,150		9/2010	Patei Kohlhase	2006/0147330		7/2006	
,	28,890			Henglein	2006/0159920			Reynders
	50,775		12/2010		2006/0225609			Rueger
	75,112		1/2011		2006/0241211			Coughlin
	59,727		6/2011		2006/0280705			Bruechert
	93,443 93,444		8/2011 8/2011		2006/0280706 2007/0028799		2/2006	Sebillotte Arnaud Kniess
	98,266			Morimitsu	2007/0028799		3/2007	
	07,583		8/2011		2007/0065381		3/2007	Elsbrock
8,0	16,934	B2	9/2011	Misaki	2007/0077218			Weinling
8,0	67,090	B2	11/2011	Domnick	2007/0134174	A1	6/2007	Irwin

US 9,168,394 B2 Page 3

(56)		Referen	ces Cited	2011/0113984 2011/0118384			Fuller, Jr. Bugnon
	U.S.	PATENT	DOCUMENTS	2011/0212042	A1	9/2011	Maitra
2007/0124177	4.1	C/2007	7:	2011/0223218 2011/0226161		9/2011	Jones Schumacher
2007/0134177 2007/0141001		6/2007	Zimmermann Clapp	2011/0236332	A1	9/2011	Dop
2007/0141002		6/2007	Montezinos	2011/0237683			Schmid
2007/0166534			Entenmann	2011/0251293 2011/0269845		10/2011	Trummer Rujard
2007/0225424 2007/0248560		9/2007 10/2007		2011/0298207			Despland
2007/0274938		11/2007	Alfano	2011/0306678		12/2011	
2007/0297996		12/2007		2012/0027862 2012/0039833			Schmidt Brennan
2008/0019933 2008/0038360			Thevenet Zukowski	2012/0039833			Shimizu
2008/0044366	A1	2/2008	Dumousseaux				
2008/0081057 2008/0102269			Chevalier Herzing	FC	REIG	N PATE	NT DOCUMENTS
2008/0102209			Thevenet	EP	696	674 D1	1/1009
2008/0127429	A1	6/2008	Brun	EP EP		674 B1 675 B1	1/1998 2/1998
2008/0168924 2008/0181921			Melson DeLuca	EP	736	073 B1	3/1998
2008/0200560		8/2008		EP EP		105 B1	9/1998
2008/0207772	A1	8/2008	Kniess	EP		512 B1 626 B2	1/1999 11/1999
2008/0210133			Roesler Birman	EP	741	170 B1	12/1999
2008/0213322 2008/0226574			Thevenet	EP EP		091 B2	1/2000
2008/0247977	A1	10/2008	Le Gendre	EP EP		343 B1 573 B1	2/2001 7/2001
2008/0249209 2008/0279796			Trummer Handrosch	EP	848	044 B1	5/2002
2008/0279790		11/2008		EP EP		959 B1 755 B1	11/2002 1/2003
2008/0292567	A1	11/2008	Schuster	EP		451 B9	2/2003
2008/0295737 2008/0314284		12/2008 12/2008	Henglein	EP	1038	941 B1	5/2003
2008/0314284		12/2008		EP EP		571 B1 698 B1	6/2003 7/2003
2009/0011035	A1		Zukowski	EP		309 B1	9/2003
2009/0013906 2009/0028808		1/2009 1/2009	Fischer	EP	1078	975 B1	10/2003
2009/0028809		1/2009		EP EP		776 B1 974 B1	12/2003 3/2004
2009/0030113			Glockner	EP		599 B1	9/2004
2009/0035241 2009/0041695		2/2009	Cassin Dumousseaux	EP		730 A1	11/2004
2009/0053164		2/2009	Opper-Linnert	EP EP		731 A1 097 B1	11/2004 1/2005
2009/0056591			Schmidt	EP	1235	882 B1	1/2005
2009/0123403 2009/0185992		5/2009 7/2009		EP EP		079 B1 821 B2	6/2005
2009/0196841	A1	8/2009	Song	EP EP		277 B1	10/2005 11/2005
2009/0208436 2009/0220557		8/2009 9/2009	Hollman Pfaff	EP		921 A1	11/2005
2009/0246294			Hochstein	EP EP		545 B2 790 B1	5/2006 8/2006
2009/0249979		10/2009		EP		535 B1	2/2007
2009/0252695 2009/0252772		10/2009 10/2009	Peng Henglein	EP		699 B1	5/2007
2009/0311209		12/2009		EP EP		864 B1 622 B1	7/2007 8/2007
2010/0011992		1/2010		EP		192 B1	2/2008
2010/0047291 2010/0047300		2/2010	Hochstein Kaupp	EP		443 B1	3/2008
2010/0089291	A1	4/2010	Kang	EP EP		906 B1 112 B1	6/2008 10/2009
2010/0095868 2010/0104610		4/2010	Kaupp Dueva-Koganov	EP	1339	375 B1	11/2009
2010/0104010		5/2010		EP EP		527 A2 665 B1	12/2010 6/2011
2010/0129412		5/2010	Kitamura	EP		586 B2	10/2011
2010/0136068 2010/0158830		6/2010 6/2010		GB	1268		7/1978
2010/0178308		7/2010		GB GB	1430 1464		9/1978 11/1978
2010/0183535		7/2010		GB	1517		3/1981
2010/0192802 2010/0196296		8/2010 8/2010	Geissler	GB	1533		1/1998
2010/0197805	A1	8/2010	Cassin	GB GB	2055 1525		5/1998 7/2005
2010/0203093		8/2010	J	JP	10194	912	3/1972
2010/0209464 2010/0218703		9/2010	Maderazzo Bujard	JP JP	10259		4/1976 2/1977
2010/0221205	A1	9/2010	Blin	JP JP	11116 0543		2/1977 2/1993
2010/0297045 2010/0322883		11/2010 12/2010	* *	JP	7309	715	11/1995
2010/0322883		12/2010		JP JP	8217 8269		8/1996 10/1996
2011/0033400	$\mathbf{A}1$	2/2011	Ehlis	JP	9012		1/1997
2011/0064779			Gruener Zhong	JP	9194		7/1997
2011/0070273 2011/0112234		3/2011 5/2011	Zneng Hall-Goulle	JP JP 2	9309 005187		12/1997 7/1998

US 9,168,394 B2

Page 4

(56)	Dofous	nces Cited	JР	2009185029	8/2009
(56)	Keiere	nces Chea	JP	2009183029	12/2009
	EODEICM DATE	ENT DOCUMENTS	JP	2009280507	12/2009
	FOREIGN PATE	ENT DOCUMENTS	JP	2010083792	4/2010
TD	10017120	0/1000	JP	2010105936	5/2010
JP	10017438	9/1998	JP	2010105930	6/2010
JP	10114867	4/1999	JP	2010120443	10/2010
JP JP	11116441 11124314	4/1999 5/1999	JP	2010235530	10/2010
JP JP	11124314	3/1999 8/1999	ĴР	2010280607	12/2010
JP JP	2000034203	8/1999 2/2000	WO	WO 9603962	2/1996
JP JP	2000034203	2/2000	WO	WO 9634917	11/1996
JP JP	2001039847	10/2001	WO	WO 9739066	10/1997
JР	2001288039	1/2002	WO	WO 9850471	11/1998
JР	2002003337	3/2002	WO	WO 0051551	9/2000
JP JP	2002087933	8/2002	WO	WO 0075240	12/2000
JP	2002241228	1/2003	WO	WO 0216505	2/2002
JP	2003003089	1/2003	WO	WO 2004007624	1/2004
JP	2003020337	6/2003	WO	WO 2004/100922	11/2004
JP	2003171232	6/2003	WO	WO 2006097352	9/2006
JP	2003171373	7/2003	WO	WO 2007/031970	3/2007
JP	2003212725	7/2003	WO	WO 2007055529	5/2007
JP	2003261423	9/2003	WO	WO 2007093334	8/2007
JР	2004010541	1/2004	WO	WO 2007118570	10/2007
JP	2004123682	4/2004	WO	WO 2007140897	12/2007
ĴР	2004238337	8/2004	WO	WO 2008007334	4/2008
JP	2004339185	12/2004	WO	WO 2008074654	6/2008
JР	2005126328	5/2005	WO	WO 2008092529	8/2008
JP	2005255633	9/2005	WO	WO 2008032323 WO 2008132042	11/2008
JP	2005314390	11/2005	wo	WO 2008135383	11/2008
JP	2005314391	11/2005	wo	WO 2008133383 WO 2009071529	6/2009
JР	2005314392	11/2005	WO	WO 2009071329 WO 2009135784	11/2009
JР	2005314394	11/2005	WO	WO 2009153784 WO 2009152907	12/2009
JР	2005314396	11/2005	WO	WO 2009132907 WO 2010050194	5/2010
JP	2005350407	12/2005			
JР	2006045562	2/2006	WO	WO 2010057968	5/2010
JР	2006328070	12/2006	WO	WO 2010063430	6/2010
JP	2007126482	5/2007	WO	WO 2010146570	12/2010
JP	2007291066	11/2007	WO	WO 2011079160	6/2011
JP	2008230997	10/2008	WO	WO 2011085780	7/2011
JP	2008255012	10/2008	WO	WO 2011095326	8/2011
JP	2009173606	8/2009	WO	WO 2012046798	4/2012

PIGMENTED SKIN-CARE COMPOSITIONS

FIELD OF THE INVENTION

The present invention relates to skin care compositions 5 and, in particular, pigmented skin care compositions that are useful in improving the visible appearance of skin.

BACKGROUND OF THE INVENTION

A variety of products are available to consumers to aid in improving skin appearance, and in particular minimizing the visibility of discontinuities in skin appearance. One way to accomplish this is through the use of compositions including colored pigments. For example, cosmetic foundations (either 1: in powdered or lotion form) contain colored pigments that are intended to mimic the skin's natural color. A problem with foundations of this type is that if such foundations are applied in a localized manner (e.g. only in a selected area of redness) the difference between the foundation color and the surround- 20 ing natural skin color may be readily apparent resulting in an irregular skin appearance. Alternatively, if such foundations are applied in a continuous manner over a larger area of skin, to thereby provide a more uniform skin appearance, an artificial "mask-like" appearance may result. It is further noted 25 acceptable carrier, at least a first and a second interference that in order to effectively conceal skin discontinuities, foundations of the type described above require relatively high pigment contents, which further contributes to the undesirable "mask-like" qualities of such compositions.

Certain colored lotions known in the art are specifically 30 intended to reduce "redness", that is reduce the red appearance of certain skin discontinuities. These redness reduction lotions typically rely upon classic pigments or dyes that appear green (the complementary color of red) to thereby deliver the redness reduction effect. The problem with such 35 by weight, unless otherwise specifically mentioned. redness reduction lotions is that they can impart an unnatural green color to the healthy skin surrounding the skin discon-

Efforts have been made to overcome the shortcomings of the products described above by way of personal care com- 40 positions that utilize interference pigments, in lieu of conventional pigments. Interference pigments typically are thin plate-like, colorless, particles including two or more layers. The layers of the interference pigment have different refractive indices, and reflect a color resulting from the constructive 45 or destructive interference of reflections of light from the different lavers.

A specific skin care composition including interference pigments is disclosed in US Patent Publication 2007065381A to Elsbrock, et al. Elsbrock discloses a skin care composition 50 that includes a first pigment that reflects a first color and a second pigment that reflects a second complementary color.

The inventors of the present invention have discovered that one drawback of the composition disclosed in Elsbrock et al. is that a formulator making a composition in accordance with 55 the teachings of Elsbrock et al. is limited in the selection of interference pigments that may be employed. Specifically, since the composition of Elsbrock et al. requires the use of a combination of a first and second pigment that reflect complementary colors, a formulator is thus limited in the selection of 60 interference pigments that may be employed. In addition, the inventors of the present invention have discovered that if redness reduction is desired, using a composition of the type disclosed in Elsbrock et al., may require the use of relatively high total pigment concentrations. Finally, the inventors of 65 the present invention have discovered that since the composition of Elsbrock et al. requires the use of a combination of a

2

first and a second pigment that are complementary colors, if the chroma of the composition is to be minimized the first and second pigments must be employed in a 1:1 ratio or substantially similar ratio.

In view of the above, the inventors have recognized that further improvements in skin care compositions including interference pigments are required. More specifically, the inventors of the present invention have recognized the need for compositions that minimize the red appearance of certain skin discontinuities with minimal impact on the appearance of the healthy skin surrounding the skin discontinuity. The inventors of present invention have also recognized the need for skin care compositions including interference pigments that can effectively reduce the red appearance of certain skin discontinuities at low total pigment concentrations. Finally, the inventors of the present invention have recognized the need for skin care compositions including at least a first and second interference pigment that have a relatively low chroma over a wide range of first and second pigment ratios.

SUMMARY OF THE INVENTION

A skin care composition including a dermatologically pigment, wherein a ratio of the first to second interference pigment is in a range between 20:80 and 80:20, wherein a chroma of said composition over said range is less than 8.5, and wherein a redness reduction index (RRI) of said composition is less than -7.5.

DETAILED DESCRIPTION OF THE INVENTION

All percentages listed in the specification are percentages

As used herein, the term "skin care" means the treatment of the human body, in certain embodiments preferably topical treatment, including, but not limited to application of composition to mammalian skin to improve the appearance of the skin, including the self-perception of one's skin. While the term "skin," is meant broadly, to include all keratinacious parts of the body (including hair and nails), in certain preferred embodiments "skin" is meant to be exclusive of hair and nails.

The present inventors have surprisingly found that it is possible to minimize the visibility of discontinuities in mammalian skin and improve the overall skin appearance by using skin care compositions according to the present invention. Skin care compositions according to the present invention include at least a first and second interference pigment. Skin care compositions according to the present invention may optionally include more than two interference pigments.

Herein, "minimize the appearance of visible discontinuities in mammalian skin" means improving the appearance of mammalian skin such that positive change in skin appearance after topically applying the composition of the present invention to the skin is observed at a distance of two feet from the user, relative to the appearance of the skin prior to application of the composition. "Visible discontinuities" include, but are not limited to, discoloration due to hyper-pigmentation, age spots, freckles, acne, scar tissue, wound, abrasion, under-eye circles, and uneven skin tone.

Applicants have also surprisingly found that it is possible to reduce the red appearance of certain skin discontinuities with minimal impact on the appearance of the healthy skin surrounding said skin discontinuity by using skin care compositions according to the present invention.

The compositions of the present invention include "interference pigments." Interference pigments typically are thin plate-like, colorless, particles including two or more layers. The layers of the interference pigment have different refractive indices, and reflect a color resulting from the constructive or destructive interference of reflections of light from the different layers. Certain interference pigments that are useful in the present invention are those that are formed from mica or borosilicate flakes coated with thin films of TiO_2 or Fe_2O_3 .

Interference pigments suitable for use in compositions of 10 the present invention have a particle size range wherein fifty percent of the particles fall within the size range (D50) of about 2 μ m and about 75 μ m. Particle size may be determined using a Malvern Mastersizer S particle size analyzer, commercially available from Malvern Instruments Ltd., Worcestershire, United Kingdom. When the term "particle size" is used herein this term is meant to refer to the D50 particle size.

Commercially available interference pigments suitable for use in the present invention are available from BASF Corporation, Florham Park, N.J. (select pigments from those sold under the Flamenco®, Lumina® and Reflecks® tradenames), from Impact Colors, Inc., Newark, Del. (select pigments from those sold under the GeminiTM tradename), and from Kobo Products, Inc., South Plainfield, N.J. (select pigments from those sold under the KTZTM tradename).

In certain embodiments of the present invention, the composition includes at least a first and second interference pigment wherein the first interference pigment has a hue angle (h°) between 180°-224° and wherein the second interference pigment has a hue angle (h°) between 45°-135°. In certain 30 embodiments of the present invention the second interference pigment has a hue angle (h°) between 80°-95°. Hue angle (h°) was measured using an X-Rite MA98 Multi-Angle Spectrophotometer, commercially available from X-Rite, Inc., Grand Rapids, Mich. To determine the hue angle (h°) of each of the 35 pigments used in the inventive compositions described in detail herein, a 3% pigment in lacquer (clear nail lacquer Sally Hansen, Hard as Nails, Nail Color 4860-01 Invisible, commercially available from Coty, Inc., New York, N.Y.) suspension was formulated. The suspension was drawn down 40 onto the black portion of a Laneta Test Opacity Chart 2A, using an Elcometer 4340 Motorized Applicator machine (the Elcometer 4340 is commercially available from Elcometer Ltd of Manchester, UK; Leneta test charts are commercially available from The Leneta Company, Mahwah, N.J.). The 45 drawdown was performed onto the test card at a speed setting of 1 with the bird applicator, resulting in a 8 mil application in a 3" wide area. After coating the test card, the card was then allowed to dry overnight prior to taking measurements. Measurements were taken of the test card using the 45 as 15 and 45 50 as 45 settings on the X-Rite MA98 Multi-Angle Spectrophotometer to measure the hue angle (h°).

In order for a pigment to be useful in compositions according to the present invention, and therefore considered within the scope of the present invention, the measured hue angle 55 (h°) at both settings must fall within the specified range of between 180° - 224° for the first interference pigment or between 45° - 135° for the second interference pigment. In certain embodiments of the invention the second interference has a hue angle (h°) between 80° - 95° . If the measured hue 60 angle (h°) for a pigment is outside the specified range at either setting (i.e. at 45 as 15 and 45 as 45) then the pigment is considered outside the scope of the present invention.

Herein, "chroma," describes color and color intensity. For the purposes of the present invention, color is defined according to a value on the CIELAB color system, which is based on the XYZ color system, defined by the Commission Interna4

tionale de l'Eclairage (CIE system) to provide a manner of objectively representing perceived color and color differences. X, Y and Z can be expressed in a variety of manners, or "scales," one of which is the Hunter scale. The Hunter scale has three variables, L, a, and b, which correlate mathematically to X, Y and Z, and is described by Robertson, A. R. in "The nCIE 1976 Color Difference Formulas," Color Research Applications, vol. 2, pp. 7-11 (1977). The compositions of the present invention may be analyzed with a Konica Minolta CR-400 Chroma Meter (commercially available from Konica Minolta Sensing Americas, Inc., Ramsey, N.J.), which generates values for L, a, and b. The value for "a" correlates to a value along the red-green (horizontal) axis, and the value for "b" correlates to a value along the blue-yellow (vertical) axis. For example, a blue-colored sample will have a negative b-value, whereas a red-colored sample will have a positive a-value. A more positive or negative value represents a more intense color. The value for "L" is an indicator of lightness and/or darkness, and correlates to a value along the z-axis, which is perpendicular to both the horizontal and vertical axes. "Chroma" is measured by a vector having its origin at the intersection of the red-green and blue-yellow axes and extending outward into the color space defined by the horizontal and vertical axes of the CIELAB color system. The length of the vector represents the chroma, and the direction of the vector represents the shade, or hue. The shorter the vector, the less colored is the composition, and the lower the chroma.

The inventors of the present invention have discovered that skin care compositions of the present invention can be effectively employed to reduce the red appearance of skin discontinuities with minimal impact on the appearance of the surrounding healthy skin. As noted above, skin care compositions according to the present invention include at least a first and second interference pigment. Surprisingly, the inventors of the present invention have discovered that the redness reduction benefit can be delivered by way of the skin care compositions of the present invention over a relatively wide range of first and second interference pigment ratios. In particular, in certain embodiments, skin care compositions of the present invention can deliver the redness reduction benefit over a first interference pigment to second interference pigment ratio range between 20:80 and 80:20.

The inventors of the present invention have discovered that the interference pigments used in the inventive compositions have unique properties when used in combination. Specifically, the interference pigments of certain embodiments of the present invention when used in combination provide a relatively low chroma over a wide range of pigment ratios. In particular, in certain embodiments of the invention, the first and second interference pigments may be employed over the entire 20:80 to 80:20 ratio range and all of such compositions will have a chroma of less than 8.5 when formulated using a substantially colorless carrier of the type described in Table 4 herein. Accordingly, when the language "wherein chroma of said composition over the entire range is less than 8.5" is used herein it means that when a plurality of different compositions are formulated using the first and second interference pigments employed in the inventive composition, and a substantially colorless carrier of the type set for in Table 4, all of such compositions will have a chroma less than 8.5. For example, if a plurality of compositions are formulated using first and second interference pigments according to certain embodiments and the carrier set forth in Table 4 at the pigment ratios of 20:80; 30:70; 40:60; 50:50; 60:40; 70:30; and 80:20 all of such compositions will have a chroma of less than 8.5. The above properties provide the formulator significantly

more flexibility than the compositions disclosed in the prior art in that the formulator can provide a relatively low chroma composition over a wide range of first to second pigment ratios

The inventors of the present invention have discovered that skin care compositions of the present invention can be effectively employed to reduce the red appearance of certain skin discontinuities with minimal impact on the appearance of the surrounding healthy skin. Surprisingly, the inventors of the present invention have discovered that this benefit can be delivered by way skin care compositions according to the present invention having relatively low total pigment concentrations. In particular, the compositions of the present invention may comprise a total amount of interference pigments of from 0.1% to 5% by weight, and in certain embodiments from 1.0% to 4%.

The inventors of the present invention have discovered that skin care compositions of the present invention can be effectively employed to reduce the red appearance of certain skin discontinuities with minimal impact on the appearance of the surrounding healthy skin. Surprisingly, the inventors of the present invention have discovered that this benefit can be delivered despite the skin care compositions not having a chroma that approaches zero. In particular, compositions according to certain embodiments of the present invention have a chroma greater than 3.5, and certain embodiments between 4.0 and 15.0.

Compositions of the present invention include a carrier useful for delivering the pigment to the human body. In certain embodiments the composition includes a cosmetically-acceptable carrier that is useful for distributing the pigment evenly across an expanse of skin. As used herein, the term "cosmetically-acceptable carrier" means a carrier that is suitable for use in contact with the skin without undue toxicity, incompatibility, instability, irritation, allergic response, and the like. This term is not intended to limit the carrier for use solely as a cosmetic (e.g., the ingredient/product can be used as a pharmaceutical).

The cosmetically-acceptable topical carrier generally 40 includes one or more of the following fluids: water, hydrophobic compounds (e.g., hydrocarbons suitable for use in cosmetic products, such as those having carbon chains from about C6 to about C50, more preferably from about C8 to about C22, such as oils, fatty esters, fatty alcohols, fatty 45 esters; as well as silicone fluids/oils) such as ones suitable to provide emolliency, spreadability, or phase stability; glycols such as ones suitable to provide humectancy such as glycerol, or diols such as propylene glycol or butylene glycol; or lower alcohols such as those suitable to provide solvency or spreadability such as ethanol or isopropanol.

In certain embodiments the carrier may include one or more solid, semi-solid, paste-like, or powder materials useful in helping to distribute the pigment across the skin. Examples include hydrophobic compounds, including waxes and other 55 hydrophobic compounds that melt above ambient temperature; powders such as starch, talc, corn starch and the like.

In a preferred embodiment, the embodiment, the cosmetically acceptable topical carrier is present in a concentration that is from about 20% to about 99.9%, preferably form about 50% to about 99.8%, more preferably from about 75% to about 99.5%. In another embodiment, the cosmetically-acceptable topical carrier includes a substrate useful for wiping the composition onto the skin. In yet another embodiment, the cosmetically-acceptable topical carrier includes a bandage 65 for applying the composition to the skin and/or maintaining the composition in contact with the skin.

6

Preferably the cosmetically acceptable topical carrier is substantially colorless. That is, the carrier does not impart any substantial color to the overall composition independent of the interference pigments employed in the composition. A suitable substantially colorless carrier is disclosed in the Table 4 although other substantially colorless carriers will be readily apparent to those of skill in the art.

The interference pigments may be distributed into the composition via techniques known to those skilled in the art. For example, the pigment may be suspended or dispersed into an oil phase and/or a water phase that is present in the composition. In order to facilitate distribution of the pigment within the composition, the composition may include a stabilizing agent (e.g., a rheology modifier, a thickening agent, a dispersing agent, or similar materials). Any of a variety of commercially available stabilizing agents which are capable of imparting the appropriate viscosity to the compositions are suitable for use in this invention. If used, the thickener may, for example, be present in an amount sufficient to raise the Brookfield viscosity of the composition to a value of between about 500 to about 10,000 centipoise. Examples of suitable thickening agents nonexclusively include: crosslinked polyacrylic acids (e.g., CARBOPOL ULTREZ 10 from Noveon, Inc. of Cleveland, Ohio); mono or diesters of 1) polyethylene glycol of formula: HO—(CH₂CH₂O)_zH, wherein z is an integer from about 3 to about 200; and 2) fatty acids containing from about 16 to about 22 carbon atoms; fatty acid esters of ethoxylated polyols; ethoxylated derivatives of mono and diesters of fatty acids and glycerine; hydroxyalkyl cellulose; alkyl cellulose; hydroxyalkyl alkyl cellulose; hydrophobically-modified alkali swellable emulsions (HASEs); hydrophobically-modified ethoxylated urethanes (HEURs); xanthan and guar gums; and mixtures thereof.

The composition may include any of various surfactants, wetting agents, or emulsifiers commonly used in personal care formulations. These materials may be ionic, non-ionic, as may be selected for their ability to provide wetting, emulsification, low irritation, foam (or lack thereof), or other desired properties. Examples of suitable surfactants, wetting agents, or emulsifiers include anionics such as surfactants include those selected from the following classes of surfactants: alkyl sulfates, alkyl ether sulfates, alkyl monoglyceryl ether sulfates, alkyl sulfonates, alkylaryl sulfonates, alkyl sulfosuccinates, alkyl ether sulfosuccinates, alkyl sulfosuccinamates, alkyl amidosulfosuccinates, alkyl carboxylates, alkyl amidoethercarboxylates, alkyl succinates, fatty acyl sarcosinates, fatty acyl amino acids, fatty acyl taurates, fatty alkyl sulfoacetates, alkyl phosphates; nonionic surfactants such as polyoxyethylene derivatives of polyol esters, wherein the polyoxyethylene derivative of polyol ester (1) is derived from (a) a fatty acid containing from about 8 to about 22, and preferably from about 10 to about 14 carbon atoms, and (b) a polyol selected from sorbitol, sorbitan, glucose, α-methyl glucoside, polyglucose having an average of about 1 to about 3 glucose residues per molecule, glycerine, pentaerythritol and mixtures thereof, (2) contains an average of from about 10 to about 120, and preferably about 20 to about 80 oxyethylene units; and (3) has an average of about 1 to about 3 fatty acid residues per mole of polyoxyethylene derivative of polyol ester; amphoterics such as amphocarboxylates such as alkylamphoacetates (mono or di); alkyl betaines; amidoalkyl betaines; amidoalkyl sultaines; amphophosphates; phosphorylated imidazolines such as phosphobetaines and pyrophosphobetaines; carboxyalkyl alkyl polyamines; alkyliminodipropionates; alkylamphoglycinates (mono or di); alkylamphoproprionates (mono or di),); N-alkyl β-aminoproprionic acids; alkylpolyamino carboxylates; and cationics

such as alkyl quaternaries (mono, di, or tri), benzyl quaternaries, ester quaternaries, ethoxylated quaternaries, alkyl amines

The compositions may include any of various other functional ingredients such as chelating agents (e.g., EDTA); pH 5 adjusters (citric acid, sodium hydroxides, and the like); preservatives; and the like.

Furthermore, compositions of the present invention may also include a skin benefit agent. A skin benefit agent is any element, an ion, a compound (e.g., a synthetic compound or a 10 compound isolated from a natural source) or other chemical moiety in solid (e.g. particulate), liquid, or gaseous state and compound that has a cosmetic or therapeutic effect on the skin. As used herein, the term "benefit agent" includes any active ingredient such as a cosmetic or pharmaceutical, that is 15 to be delivered into and/or onto the skin, hair, mucosa, or teeth at a desired location.

Examples of suitable benefit agents include those that provide benefits such as, but not limited to: de-pigmentation agents; amino acids and their derivatives; antimicrobial 20 agents; allergy inhibitors; anti-acne agents; anti-aging agents including anti-wrinkling agents and benefit agents suitable for treating loss of skin elasticity, uneven skin, blotchiness, and skin tone; tropoelasin promoters and tropoelastin crosslinkers; antiseptics; analgesics; antipruritics; local anes-25 thetics; anti-hair loss agents; hair growth promoting agents; hair growth inhibitor agents, antihistamines; antiinfectives; anti-inflammatory agents; anticholinergics; vasoconstrictors; vasodilators; wound healing promoters; peptides, polypeptides and proteins; deodorants and anti-perspirants; medicament agents; skin firming agents, vitamins; skin lightening agents; skin darkening agents; antifungals; depilating agents; counterirritants; enzymes for exfoliation or other functional benefits; enzyme inhibitors; NFkB-inhibitors; herbal extracts; flavenoids; sensates and stress-reducing agents; 35 anti-oxidants; hair lighteners; sunscreens; anti-edema agents, neo-collagen enhancers, anti-dandruff/sebhorreic dermatitis/ psoriasis agents; keratolytics; and mixtures thereof.

Particularly suitable benefit agents include depigmentation agents and skin-lightening agents such as hydroquinone and 40 extracts of soy; keratolytic agents and/or anti-acne agents such as alpha and beta hydroxyacids such as salicylic acid; anti-aging actives such as retinoids including retinol, amines such as N,N,N',N'-Tetrakis(2-hydroxypropyl)ethylenediamine (THPED), N,N, N',N'-Tetrakis(2-hydroxyethyl) ethylenediamine (TEMED), substituted resorcinols such as 4-hexyl resorcinol, anti-inflammatories such as extracts of feverfew, tropoelastin promoters such as extracts of blackberry, tropoelastin crosslinkers such as extracts of dill, among other 50 skin benefit agents.

The amount of the benefit agent that may be used may vary depending upon, for example, the ability of the benefit agent to penetrate through the skin, nail, mucosa, or teeth; the specific benefit agent chosen, the particular benefit desired, 55 the sensitivity of the user to the benefit agent, the health condition, age, and skin and/or nail condition of the user, and the like. In sum, the benefit agent is used in a "safe and effective amount," which is an amount that is high enough to deliver a desired skin or nail benefit or to modify a certain 60 condition to be treated, but is low enough to avoid serious side effects, at a reasonable risk to benefit ratio within the scope of sound medical judgment.

The compositions may be made into a wide variety of product types that include but are not limited to cleansing 65 liquid washes, gels, sticks, sprays, solid bars, shampoos, pastes, foams, powders, mousses, wipes, patches, hydrogels,

8

and films. These product types may comprise several types of cosmetically-acceptable carriers including, but not limited to solutions, emulsions (including for example, oil-in-water, water-in-oil, microemulsions and nanoemulsions, and the like), gels, and solids. The following are non-limitative examples of such carriers. Other carriers can be formulated by those of ordinary skill in the art.

The compositions useful in the present invention can be formulated as solutions. Solutions typically include an aqueous or organic solvent (e.g., from about 50% to about 99.99% or from about 90% to about 99% of a cosmetically acceptable aqueous or organic solvent). Examples of suitable organic solvents include: polyglycerols, propylene glycol, polyethylene glycol (200, 600), polypropylene glycol (425, 2025), glycerol, 1,2,4-butanetriol, sorbitol esters, 1,2,6-hexanetriol, ethanol, and mixtures thereof. In certain preferred embodiments, the compositions of the present invention are aqueous solutions comprising from about 50% to about 99% by weight of water.

According to certain embodiments, compositions useful in the subject invention may be formulated as a solution comprising an emollient. Such compositions preferably contain from about 2% to about 50% of an emollient(s). As used herein, "emollients" refer to materials used for the prevention or relief of dryness, as well as for the protection of the skin. A wide variety of suitable emollients are known and may be used herein. Sagarin, Cosmetics, Science and Technology, 2nd Edition, Vol. 1, pp. 32 43 (1972) and the International Cosmetic Ingredient Dictionary and Handbook, eds. Wenninger and McEwen, pp. 1656 61, 1626, and 1654 55 (The Cosmetic, Toiletry, and Fragrance Assoc., Washington, D.C., 7.sup.th Edition, 1997) (hereinafter "ICI Handbook") contains numerous examples of suitable materials. A lotion can be made from such a solution. Lotions typically comprise from about 1% to about 20% (e.g., from about 5% to about 10%) of an emollient(s) and from about 50% to about 90% (e.g., from about 60% to about 80%) of water.

The present compositions may be of varying phase compositions, including those having an exterior aqueous phase (e.g., aqueous phase is the most exterior phase of the composition). As such, compositions of the present invention may be formulated to be oil-in-water emulsions that are shelf-stable in that the emulsion does not lose phase stability or "break" when kept at standard conditions (22 degrees Celsius, 50% relative humidity) for a week or more after it is made.

For those compositions that include an aqueous phase, the pH of the present compositions is not critical, but may be in a range that does not facilitate irritation to the skin, such as from about 4 to about 7. The viscosity of the personal care composition is not critical, although it may be a spreadable cream or lotion or gel.

The pigment, carrier and optional other components of the composition may be combined according to the present invention via any conventional methods of combining two or more fluids or solids. For example, one or more compositions comprising, consisting essentially of, or consisting of at least one pigment and one or more compositions comprising, consisting essentially of, or consisting of water or suitable ingredients may be combined by pouring, mixing, adding dropwise, pipetting, pumping, and the like, one of the compositions comprising the polymerized surfactant into or with the other in any order using any conventional equipment such as a mechanically stirred propeller, paddle, and the like.

In certain embodiments, the composition may be impregnated within a substrate (e.g., non-woven fibrous material, a film material, or combinations thereof). The substrate material may be selected to facilitate depositing the pigment on the

skin. The methods of the present invention may further comprise any of a variety of steps for mixing or introducing one or more of the optional components described hereinabove with or into a composition comprising the pigment, either before, after, or simultaneously with the combining step described 5

In certain embodiments, the compositions produced via the present invention are preferably used as or in personal care products for treating at least a portion of a mammalian body, for example, the human body. Examples of certain preferred personal care products include various products suitable for application to the skin or hair. Particularly preferred products are those that are designed to be applied to the skin and not immediately rinsed off. Examples of these "leave-on" products particularly for use on the face, but also including those for the body, hands, feet, and the like.

As discussed above, the inventors of the present invention have surprisingly found that it is possible to reduce the red appearance of skin discontinuities with minimal impact on the appearance of the healthy skin surrounding said skin discontinuity by using skin care compositions according to the present invention. The Redness Reduction Index (RRI) Test set forth in detail below was used to illustrate the inventive skin care compositions ability to reduce the red appearance of skin discontinuities. The Healthy Skin Color Change Value (HSCCV) Test set forth in detail below was used to illustrate inventive skin care compositions ability to deliver redness reduction with minimal impact on the appearance of the healthy skin surrounding the skin discontinuity. The $_{30}$ inventors of the present invention have further discovered that in order for a skin care composition to preserve the natural tone of healthy skin it cannot impart a "green" tone to the healthy skin. Furthermore, any such green tone must not be visible when the skin is viewed from multiple angles since the human eye simultaneously views a surface from multiple angles. The Multi-Angle Δa* (MADA) Test set forth below was used to illustrate the inventive skin care compositions ability to preserve the natural tone of healthy skin without multiple angles.

Redness Reduction Index (RRI) Test

A test card including a first portion that is representative of a healthy Caucasian skin color and a second portion that is representative of an inflamed skin discontinuity was prepared as follows. The test card had dimensions of 4"x6" and was formed from Olympus P100 photo paper. The photo paper was printed using an Olympus P-10 printer which was loaded with the bundled ink ribbon. The first and second portions, each measuring 1.3"×6", were created by entering the RGB values, set forth in Table 1 below, into MATLAB R2011a software (commercially available from Mathworks, Inc., Natick, Mass.) and then printing the test card.

TABLE 1

	R	G	В
Healthy Skin	238	203	181
Inflamed Skin	170	53	65

A mimic stratum corneum overlay was prepared as follows. A stratum corneum mimic layer formed from VITRO-CORNEUM® (commercially available from IMS, Inc., Portland, Me.) was attached in a non-hydrated state to a cellulose acetate slide measuring 5"x7" (PP2500 Transparency Film 65 for Plain Paper Copiers, commercially available from 3M, St. Paul, Minn.) using collagen glue (Resine ou Pigment Pur,

concentration 8% in water, commercially available from Sennlier, Paris, France) such that it covered the entire slide, with the rough surface of the VITRO-CORNEUM® facing outward. After setting, the prepared substrate was cut into strips measuring 1.5 cm×6.0 cm, each strip constituting one mimic stratum corneum overlay.

After printing of the test card, the color of the test card was assessed using a Hunter UltraMax Colorimeter (commercially available from Hunter Associates Laboratory, Inc., Reston, Va.), both with and without the mimic stratum corneum overlay. The readings for the test card alone are set forth in Table 2 below and the readings for the test card measured through the mimic stratum corneum overlay (i.e. with the stratum corneum overlay on top of the test card, with the VITRO-CORNEUM® surface facing towards the measurement device) are set forth in Table 3 below.

TABLE 2

_	Test Card				
	L*	a*	b*		
Healthy Skin	84.28	9.71	16.05		
Inflamed Skin	40.71	48.60	21.08		

TABLE 3

	_	est Card with Mi tum Corneum O	
	L*	a*	b*
Healthy Skin	81.45	9.21	16.54
Inflamed Skin	48.86	22.22	10.61

The pigmented skin care composition to be assessed was imparting a "green" tone thereto when the skin is viewed from 40 finger applied on the mimic stratum corneum overlay in an amount of 2.5 mg/cm² and smoothed until a consistent layer was attained and then allowed to dry at room temperature overnight. To assess the pigmented skin care composition's impact on the inflamed and healthy skin color targets, the treated mimic stratum corneum overlay was placed over the two color targets on the test card and using a Hunter UltraMax colorimeter. L*, a* and b* values were measured over both portions of the test card, that is separate L*, a* and b* measurements were conducted over the "Healthy Skin" portion of the test card through the treated mimic stratum corneum overlay and separately over the "Inflamed Skin" portion of the test card through the treated mimic stratum corneum

> Using the measured values for a* over the "Inflamed Skin" 55 portion, and the original a* value of the test card over the "Inflamed Skin" portion of the test card set forth in Table 3 above, Δa* was calculated according to the equation $\Delta a^*=a^*_2-a^*_1$, where a^*_2 is the a^* value measured through the treated mimic stratum corneum overlay over the "Inflamed Skin" portion of the test card and a*1 is the original a* value set forth in Table 3 above for the "Inflamed Skin" portion of the test card measured through the non-treated mimic stratum corneum overlay. This measured value of Δa^* indicates the ability of the test composition to reduce the "red" appearance of a skin discontinuity, the more "negative" the Δa^* value the greater the redness reduction capabilities of the test composition.

The above described test was repeated four (n=4) times for each test composition. An average was taken of each of the four calculated Δa^* values to provide an average Δa^* value. This average Δa* is referred to herein as the Redness Reduction Index (RRI).

Compositions according to the present invention have a redness reduction index (RRI) of less than -7, in some embodiments less than -7.5, in some embodiments less than -9.5, in some embodiments between -7.5 and -16.0, and in some embodiments between -10.0 and -16.0.

Healthy Skin Color Change Value (HSCCV) Test

Using the values for L*, a* and b* measured through the treated mimic stratum corneum overlay over the "Healthy Skin" portion of the test card, and the original L*, a* and b* values of the "Healthy Skin" of the test card measured through the non-treated mimic over the "Healthy Skin" portion of the test card (set forth in Table 3 above), the ΔE of the "Healthy Skin" portion of the test card was determined according to the formula set forth below.

where

 L_2^* , a_2^* , and $b_2^*=L^*$, a^* and b^* value of test card in the healthy skin portion of the card as measured through the 25 treated mimic stratum corneum layer; and

 L_1^* , a_1^* and $b_1^*=L_1^*$, a_1^* and a_1^* value of test card in the healthy skin portion of the card as measured through the non-treated mimic stratum corneum layer (set forth in Table 3

The calculated ΔE value indicates the degree to which the test composition changed the appearance of healthy skin, the smaller the ΔE value the less the test composition altered the appearance of healthy skin.

The above described test was repeated four (n=4) times for 35 each test composition. An average was taken of each of the four calculated ΔE values to provide an average ΔE value. This average ΔE is referred to herein as the Healthy Skin Color Change Value (HSCCV). Compositions according to the present invention have an HSCCV of less than 3 and in 40 certain embodiments between 0 and 2.5. In this regard, it is noted that the human eye can barely detect a ΔE≈2.3. (Gaurav Sharma (2003). Digital Color Imaging Handbook (1.7.2 ed.). CRC Press. ISBN 0-8493-0900-X)

Compositions of the present invention simultaneously pro- 45 vide a negative RRI and a small HSCCV. The combination of a negative RRI and a small HSCCV indicates that the inventive compositions are both effective at reducing the red appearance of skin discontinuities while at the same time have minimal impact on the appearance of the surrounding 50 where healthy skin.

Chroma Measurement Test

For each of the test compositions tested above, the chroma of such composition was determined as follows.

Three grams (3 g) of the test composition was drawn down 55 onto an AF4300 Write-On Transparency Film (commercially available from 3M, St. Paul, Minn.). The drawdown was conducted by hand using the 6 mil side of a 2-path, 5 inch applicator, part #663479, Precision Gage and Tools Co., Dayton, Ohio. After coating the transparency was allowed to dry 60 overnight prior to taking measurements. After drying, the transparency was placed over the black portion of Laneta Test Opacity Chart 2A (commercially available from The Leneta Company, Mahwah, N.J.).

Using a Konica Minolta CR-400 Chroma Meter (commer- 65 cially available from Konica Minolta Sensing Americas, Inc., Ramsey, N.J.), L*, a* and b* values were measured over the

12

black portion of the test card. Using these values, chroma for the test composition was calculated according to the following equation.

Chroma=
$$\sqrt{(a^{*2}+b^{*2})}$$

The above described test was repeated three (n=3) times for each test composition. An average Chroma value was taken from the four calculated Chroma values to provide an average Chroma value. Compositions according to certain embodiments of the present invention have an average chroma greater than 3.5, and in certain embodiments between 4.0 and 15.0.

Multi-Angle Δa*Test (MADA)

For each of the test compositions tested above, the MADA of such composition was determined as follows. Using the same test card and mimic stratum corneum overlay described above in the Redness Reduction Index (RRI) Test, an a* reading was measured for the Healthy Skin portion of the test $_{20}$ card through the mimic stratum corneum overlay using an X-Rite MA98 Multi-Angle Spectrophotometer, commercially available from X-Rite, Inc., Grand Rapids, Mich. It is noted that for purposes of the MADA test the mimic stratum corneum was cut to have dimensions of 3.81 cm×3.81 cm. The a* reading was measured at eight different angles and each of these values was averaged to provide an average a* reading for the Healthy Skin portion of the test card through the mimic stratum corneum overlay. The eight different angles comprised the following angle settings on the test apparatus 45 as-15, 45 as15, 45 as25, 45 as45, 45 as75, 45 as110, 15 as-15, and 15 as15.

The pigmented skin care composition to be assessed was finger applied on the mimic stratum corneum overlay in an amount of 2.5 mg/cm² and smoothed until a consistent layer was attained and then allowed to dry at room temperature overnight.

An a* reading was measured for the Healthy Skin portion of the test card through the test composition. The a* reading was measured at eight different angles and each of these values was averaged to provide an average a* reading for the Healthy Skin portion of the test card measured through test composition. The eight different angles comprised the following angle settings on the test apparatus 45 as-15, 45 as15, 45 as 25, 45 as 45, 45 as 75, 45 as 110, 15 as -15, and 15 as 15.

Using the measured a* values an average Δa * value, or MADA value, was calculated according to the following formula:

$$\Delta a^{*}=a^{*}_{2}-a^{*}_{1};$$

a*₂=average a* value for the Healthy Skin portion of the test card measured through test composition applied to mimic stratum corneum overlay; and a*1=average a* value for the Healthy Skin portion of the test card measured through the non-treated mimic stratum corneum overlay.

The above described test was repeated four (n=4) times for each test composition. An average MADA value was taken from the four calculated MADA values to provide an average MADA value. Compositions according to the present invention have an average MADA value of greater than -4.0, and in certain embodiments between -3.5 and 0.3.

INVENTIVE EXAMPLES

Pigmented skin care compositions according to the present invention were formulated to include an oil in water cream carrier and the selected interference pigments. The amount of

water in the formulation was adjusted according to the amount of pigment employed in the specific inventive example. The base oil in water cream carrier was selected due to its low optical impact to the L*a*b* measurements conducted according to the test methods set forth above herein.

The inventive formulations are summarized in Table 4 below. The pigments employed in each inventive formulation are summarized in Table 5 below. The hue angle (h°) as measured at 45 as 15 and 45 as 45 is specified for each pigment used in each of the inventive compositions.

14

TABLE 4

Components		_
Trade Name	INCI Name	% wt/wt
	Water phase	
PURIFIED WATER	WATER	79.95-83.45
Emery 917 Carbomer 20% SODIUM HYDROXIDE	Glycerin Ultrez 10 SODIUM HYDROXIDE	5.00 0.60 Trace (to achieve pH target)
solution	Oil phase	
VERSENE NA Brij 72 Brij 721 Finsolv TN DC Q7-9120 Silicone Fluid, 20 est Phenonip XB	Disodium EDTA Steareth-2 Steareth-21 C12-15 Alkyl Benzoate Dimethicone Phenoxyethanol and methyl and ethyl and	0.20 0.75 1.50 2.00 5.00
	propyl parabens Pigments	0.50.400
		0.50-4.00
	PURIFIED WATER Emery 917 Carbomer 20% SODIUM HYDROXIDE solution VERSENE NA Brij 72 Brij 721 Finsolv TN DC Q7-9120 Silicone Fluid, 20 cst	PURIFIED WATER Emery 917 Carbomer 20% SODIUM HYDROXIDE solution VERSENE NA Brij 72 Brij 721 Finsolv TN DC Q7-9120 Silicone Fluid, 20 cst Phenonip XB WATER WATER Glycerin Ultrez 10 SODIUM HYDROXIDE HYDROXIDE SODIUM HYDROXIDE HYDROXIDE SODIUM HYDROXIDE LTTE 10 SODIUM HYDR

TABLE 5

	Pigment Composition
Inventive	Flamenco Summit Turquoise, 0.4% + Flamenco Summit Gold, 1.6%
Example #1	(45as15-189°, 45as45-193°), (45as15-88°, 45as45-84°)
Inventive	Flamenco Summit Turquoise, 0.8% + Flamenco Summit Gold, 1.2%
Example #2	(45as15-189°, 45as45-193°), (45as15-88°, 45as45-84°)
Inventive	Flamenco Summit Turquoise, 1.2% + Flamenco Summit Gold, 0.8%
Example #3	(45as15-189°, 45as45-193°), (45as15-88°, 45as45-84°)
Inventive	Flamenco Summit Turquoise, 1.6% + Flamenco Summit Gold, 0.4%
Example #4	(45as15-189°, 45as45-193°), (45as15-88°, 45as45-84°)
Inventive	Flamenco Summit Turquoise, 1% + Flamenco Summit Gold, 3%
Example #5	(45as15-189°, 45as45-193°), (45as15-88°, 45as45-84°)
Inventive	Flamenco Summit Turquoise, 1% + Flamenco Summit Gold, 2%
Example #6	(45as15-189°, 45as45-193°), (45as15-88°, 45as45-84°)
Inventive	Flamenco Summit Turquoise, 1% + Flamenco Summit Gold, 1%
Example #7	(45as15-189°, 45as45-193°), (45as15-88°, 45as45-84°)
Inventive	Flamenco Summit Turquoise, 1.32% + Flamenco Summit Gold, 0.66%
Example #8	(45as15-189°, 45as45-193°), (45as15-88°, 45as45-84°)
Inventive	Flamenco Summit Turquoise, 0.5% + Flamenco Summit Gold, 0.5%
Example #9	(45as15-189°, 45as45-193°), (45as15-88°, 45as45-84°)
Inventive	Flamenco Summit Turquoise, 1% + KTZ Interfine Gold, 1%
Example #10	(45as15-189°, 45as45-193°), (45as15-91°, 45as45-84°)
Inventive	Flamenco Summit Turquoise, 1% + KTZ Interfine Gold, 2%
Example #11	(45as15-189°, 45as45-193°), (45as15-91°, 45as45-84°)
Inventive	Flamenco Summit Turquoise, 1% + Timiron Splendid Gold, 1%
Example #12	(45as15-189°, 45as45-193°), (45as15-91, 45as45-88°)
Inventive	Flamenco Summit Turquoise, 1% + Timiron Splendid Gold, 2%
Example #13	(45as15-189°, 45as45-193°), (45as15-91, 45as45-88°)
Inventive	Lumina Turquoise 9T30D, 1% + Flamenco Summit Gold, 1%
Example #14	(45as15-193°, 45as45-207°), (45as15-88°, 45as45-84°)
Inventive	Lumina Turquoise 9T30D, 1% + Flamenco Summit Gold, 2%
Example #15	(45as15-193°, 45as45-207°), (45as15-88°, 45as45-84°)
Inventive	Reflecks MultiDimensions Transforming Teal, 1% + Flamenco Summit Gold, 1%
Example #16	(45as15-193°, 45as45-207°), (45as15-88°, 45as45-84°)
Inventive	Gemini GB-38, 1% + Flamenco Summit Gold, 1%
Example #17	(45as15-210°, 45as45-198°), (45as15-88°, 45as45-84°)

13

16

	Pigment Composition
Inventive Example #18	Gemini GB-38, 1% + Flamenco Summit Gold, 2% (45as15-210°, 45as45-198°), (45as15-88°, 45as45-84°)

In a clean beaker, combine all oil phase ingredients then began agitation and heat to 55-60° C. until the oil phase is homogeneous. In a separate clean beaker, combine water and other water phase ingredients and began agitation and heat to 55-60° C. until the water phase is homogeneous. Add the oil phase to water phase with increased agitation, mixing at a high speed for 8 minutes at 55-60° C. Cool the mixture to 50° C. and then add the Dimethicone. At 40° C., add the Phenonip, and continue mixing until uniform. Continue cooling until a temperature of 30° C. is reached, check the pH and adjust with Sodium Hydroxide solution to a target pH of 5.4 or in a range from 5.2 to 5.7

Disperse the pigment in thirty percent of the deionized water to be contained in the final composition to suspend the particles and mix thoroughly with propeller blade (or spatula depending on batch size) inside a beaker. Combine pigment premix with oil in water cream carrier prepared above in main ² beaker and mix thoroughly until uniform.

COMPARATIVE EXAMPLES

A total of eight commercial skin care products were evaluated using the test methods described above. The eight commercial products evaluated are listed below.

Eucerin Redness Relief Daily Perfecting Lotion

Comparative Example #2

Dermalogica Sheer Tint Redness Relief

Comparative Example #3

Clearasil ULTRA Overnight Face Lotion

Comparative Example #4

Eucerin Redness Relief Soothing Anti-Aging Serum

Comparative Example #5

Neutrogena Oil Free Acne Stress Control

Comparative Example #6

Clinique Redness Solutions Daily Relief Cream

Comparative Example #7

Estee Lauder Idealist Even Skintone Illuminator

Comparative Example #8

Clinique Redness Solutions Urgent Relief Cream Additional comparative examples were formulated using the 65 oil in water carrier described above in Table 4 in combination with the pigments set forth below in Table 6. TABLE 6

Comparative	KTZ Interfine ™	2.5%
Example 9	Blue KTZ Interfine ™	2.5%
Comparative Example 10	Gold KTZ Interfine ™ Red	1.0%
	KTZ Interfine ™ Green	1.0%
Comparative Example 11	KTZ Interfine ™ Gold	0.3%
•	KTZ Interfine ™ Violet	0.3%
Comparative Example 12	KTZ Interfine ™ Green	0.83%
	KTZ Interfine ™ Violet	0.34%
	Prestige Silk ™ Orange	0.83%
Comparative Example 13	KTZ Interfine TM Blue	0.5%
	KTZ Interfine TM Gold KTZ Interfine TM	0.5% 0.5%
	KTZ Interfine ™ Red KTZ Interfine ™	0.5%
Comparative	Green Timiron Splendid TM	1.0%
Example 14	Gold Timiron Splendid TM	1.0%
Comparative	Blue Timiron Splendid TM	0.4%
Example 15	Gold Timiron Splendid ™	0.35%
	Blue KTZ Interval ™	0.4%
	Red KTZ Interval ™	0.35%
Comparative	Green Prestige Silk™	1.5%
Example 16	Blue Prestige Silk™	1.5%
Comparative	Gold KTZ Interval ™	0.9%
Example 17	Red KTZ Interval ™ Green	0.8%
Comparative Example 18	KTZ Interval ™ Gold-11S2	0.75%
Zmanple 10	KTZ Interval ™ Blue-11S2	0.75%
Comparative Example 19	KTZ Interval ™ Gold-11S2	0.3%
	KTZ Interval ™ Blue-11S2	0.3%
	KTZ Interval ™ Red-11S2	0.3%
	KTZ Interval TM Green-11S2	0.3%
Comparative Example 20	Timiron Super Green	4%
Comparative Example 21 11850-164-1	Timiron Super Green	3%

The Inventive Examples and Comparative Examples described above were tested according to the Redness Reduction Index (RRI) Test, Healthy Skin Color Change Value (HSCCV) Test, Chroma Measurement Test and the Multi-Angle Δa^* (MADA) Test set forth herein. The results of such test are summarized in the Table of Results set forth below.

TABLE OF RESULTS				
SAMPLE	RRI	HSCCV	Chroma	MADA
Inventive Example 1	-12.3	2.4	7.3	-1.1
Inventive Example 2	-13.2	2.3	6.1	-1.5
Inventive Example 3	-10.9	1.8	4.8	-2.8
Inventive Example 4	-9.9	1.6	5.5	-3.8
Inventive Example 5	-17.3	3.6	14.3	-2.4
Inventive Example 6	-13.6	2.4	8.1	-1.7
Inventive Example 7	-10.8	1.7	4.9	-1.1
Inventive Example 8	-9.9	1.4	5.3	-3.5
Inventive Example 9	-7.2	1.0	3.7	-2.0
Inventive Example 10	-6.3	1.3	7.1	-2.1
Inventive Example 11	-7.7	1.5	12.3	-1.7
Inventive Example 12	-9.0	1.4	5.4	-1.2
Inventive Example 13	-12.4	2.1	6.1	-2.7
Inventive Example 14	-9.0	1.4	4.1	-3.4
Inventive Example 15	-15.4	2.7	8.1	-3.1
Inventive Example 16	-9.7	1.3	3.6	0.2
Inventive Example 17	-8.6	1.5	5.5	-1.5
Inventive Example 18	-12.9	2.4	8.1	-1.0
Comparative Example 1	-24.1	7.0	7.3	-6.9
Comparative Example 2	-7.6	3.2	7.6	-2.3
Comparative Example 3	0.0	0.6	0.2	0.8
Comparative Example 4	-1.3	2.1	1.1	-0.1
Comparative Example 5	-0.9	1.2	0.8	0.8
Comparative Example 6	-1.8	0.5	1.2	0.8
Comparative Example 7	-6.8	0.6	4.5	0.7
Comparative Example 8	-0.8	0.4	0.8	0.6
Comparative Example 9	-14.1	3.5	2.1	-0.9
Comparative Example 10	-6.9	0.9	1.3	0.1
Comparative Example 11	-0.9	0.4	1.3	1.5
Comparative Example 12	-6.8	0.9	1.0	0.3
Comparative Example 13	-6.0	0.9	1.2	0.2
Comparative Example 14	-5.6	1.1	3.3	0.4
Comparative Example 15	-5.1	0.6	0.9	0.4
Comparative Example 16	-9.2	2.4	3.5	-0.1
Comparative Example 17	-5.0	0.7	0.8	0.5
Comparative Example 18	-5.0	0.6	1.3	0.5
Comparative Example 19	-4.9	0.8	1.1	0.7
Comparative Example 20	-20.8	3.5	10.4	-5.1
Comparative Example 21	-15.4	2.0	8.7	-4.4

While particular embodiments of the present have been illustrated and described, it would be obvious to those skilled in the art that various other changes and modifications can be 40 made without department from the spirit and scope of the invention. It is therefore intended to cover in the appended claims all such changes and modifications that are within the scope of this invention.

We claim:

A skin care composition comprising:
 a dermatologically acceptable carrier;
 at least a first and a second interference pigment;
 wherein the ratio of the first to second interference pigment is in a range between 20:80 and 80:20 by weight;

- wherein the chroma of said composition over said range is less than 8.5;
- wherein the redness reduction index (RRI) of said composition is less than -7.5; and
- wherein the first interference pigment has a hue angle (h°) between 185°-215° and the second interference pigment has a hue angle (h°) between 80°-95°.
- 2. The skin care composition according to claim 1, wherein the chroma of said composition is between 4.0 and 8.0.
 - 3. The skin care composition according to claim 2, wherein the composition has a redness reduction index (RRI) of less than -9.5.
- 5 4. The skin care composition according to claim 3, wherein the composition has a redness reduction index (RRI) between -10.0 and -16.0.
- 5. The skin care composition according to claim 1, wherein the total amount of interference pigments is from 0.1% to 5% by weight.
 - **6**. The skin care composition according to claim **5**, wherein the total amount of interference pigments is from 1% to 4% by weight.
 - 7. The skin care composition according to claim 6, wherein the composition has a healthy skin color change value (HSCCV) of less than 3.
 - **8**. The skin care composition according to claim **7**, wherein the composition has a HSCCV of between 0.5 and 2.5.
 - The skin care composition according to claim 1, wherein the skin care composition additionally comprises at least one skin care active.
- 10. The skin care composition according to claim 9, wherein the skin care active is selected from the from the group consisting of depigmentation agents and skin-lightening agents, anti-acne agents anti-aging actives, and anti-inflammatories.
 - 11. The skin care composition according to claim 8, wherein each of the first and second interference pigment has a particle size within the range of 2 μ m and 75 μ m.
- 12. The skin care composition according to claim 11, wherein the composition has a multi-angle Δa^* (0) value of 45 greater than -4.0.
 - 13. The skin care composition according to claim 12, wherein the composition has a MADA value of between -3.5 and 0.3.

* * * * *